CLAIMS

 A method for producing a compound represented by formula (I):

$$R_1$$
 R_2
 R_2
 R_3
 R_3
 R_3
 R_4
 R_5
 R_5
 R_6
 R_7
 R_8

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(wherein A represents - (CH_2) n-, where n represents an integer of 0 to 10;

B represents $-CH_2-$, -(C=O)-, -CH(OH)-, $-CH(NH_2)-$ or -C(=NOR)-, where R represents a hydrogen atom, a linear or branched alkyl group having 1 to 8 carbon atoms (which may be substituted with an amino group that may be mono- or disubstituted with a linear or branched alkyl group having 1 to 4 carbon atoms);

D represents - (CH₂)_m-R', where m represents an

integer of 0 to 10, and R' represents a hydrogen atom, a

linear or branched alkyl group, a linear or branched

alkynyl group, a linear or branched alkenyl group, a

cycloalkyl group, a cycloalkenyl group, a heterocyclyl

group which may be substituted, an aryl group which may be

substituted, a heteroaryl group which may be substituted,

an -OX group (where X represents a hydrogen atom, a linear

or branched alkyl group, a linear or branched alkynyl group,

a linear or branched alkenyl group, a cycloalkyl group or

an aryl group which may be substituted) or a halogen atom;

E represents a hydrogen atom or a linear or branched

alkyl group;

G represents $-(CH_2)_p-J$, where p represents an integer of 0 to 4, and J represents a hydrogen atom, an OH group, a SH group, a methylthio group, a carboxyl group, a carbamoyl group, an amino group, a guanidino group, a linear or branched alkyl group, a cycloalkyl group, a linear or branched alkynyl group, a linear or branched alkenyl group, an aryl group which may be substituted, a heterocyclyl group which may be substituted, or a heteroaryl group which may be substituted;

bond Q represents a single bond or a double bond; and

 R_1 , R_2 and R_3 may be the same or different, and each represent a hydroxyl group, an amino group (which may be mono- or di-substituted with a linear or branched alkyl group having 1 to 4 carbon atoms), -OL, a linear or branched alkyl group, a linear or branched alkenyl group or a linear or branched alkynyl group, where L represents a linear or branched alkyl group, a linear or branched alkenyl group or a linear or branched alkynyl group), a prodrug thereof or a pharmaceutically acceptable salt thereof;

comprising reacting a compound as the starting compound represented by the following formula:

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(wherein A, D and bond Q have the same meanings as defined above, and X and Y may be the same or different and each represent a linear or branched alkyl group or a

protecting group of a carboxyl group) with an α -amino acid ester represented by the following formula:

(wherein E and G have the same meanings as defined above, and Z represents of a linear or branched alkyl group or a protecting group of a carboxyl group) in the presence of a base and a coupling agent, to yield a compound represented by the following formula:

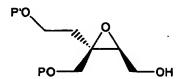
(wherein A, D, E, G, bond Q, X, Y and Z have the same meanings as defined above), and then subjecting this compound to hydrolysis, reduction, amination or amidation, hydroxyimination and/or ester conversion, if desired, to obtain the desired compound of the formula (I).

15 2. A method for producing a compound represented by the following formula:

(wherein D and n have the same meanings as defined

in claim 1, M_1 and M_2 may be the same or different and each represent an oxygen atom or a sulfur atom, and P and P' may be the same or different and each represent a hydroxyl protecting group);

comprising reacting a compound represented by the following formula:



(wherein P and P' have the same meanings as defined above) with a compound represented by the following formula:

(wherein D, n, M_1 and M_2 have the same meanings as defined above).

3. A compound represented by formula (I):

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(wherein A, B, D, E, G, bond Q, R_1 , R_2 and R_3 have the same meanings as defined in claim 1), a prodrug thereof or a pharmaceutically acceptable salt thereof.

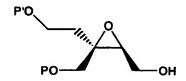
The compound of the formula (I) according to claim 3, a
 prodrug thereof or a pharmaceutically acceptable salt thereof, wherein in the case n represents 6, D represents a

n-heptyl group and p represents 1, then J represents a group which is neither a phenyl group (the phenyl group is substituted with an -OW group at the p-position where W represents a hydrogen atom, a linear or branched alkyl group, or a linear or branched alkenyl group) nor a 3-indolyl group.

- 5. The compound of the formula (I) according to claim 3, a prodrug thereof or a pharmaceutically acceptable salt thereof, wherein in the case n represents 6, D represents a n-heptyl group and p represents 1, then J represents a group which is neither a phenyl group (the phenyl group is substituted with an -OW group at the p-position where W represents a hydrogen atom, a linear or branched alkyl group, a linear or branched alkenyl group or a linear or branched alkynyl group) nor a 3-indolyl group.
- 6. A compound represented by the following formula:

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(wherein P and P' may be the same or different and each represent a hydroxyl protecting group).

20 7. A compound represented by the following formula:

(wherein A, D, X and Y have the same meanings as defined in claim 1).

A pharmaceutical composition containing a compound of
 the formula (I) according to anyone of claims 3 to 5, a
 prodrug thereof or a pharmaceutically acceptable salt

thereof.

- 9. The pharmaceutical composition according to claim 8 for preventing or treating a viral infectious disease.
- 10. The pharmaceutical composition according to claim 9 wherein the viral infectious disease is an infectious disease by HCV.
 - 11. The pharmaceutical composition according to claim 10, wherein the infectious disease by HCV is hepatitis C, cirrhosis, liver fibrosis or liver cancer.

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